Neutralizing antibody and CD8+ T cell responses following BA.4/5 bivalent COVID-19 booster vaccination in adults with and without prior exposure to SARS-CoV-2

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Neutralizing antibody and CD8⁺ T cell responses following BA.4/5 bivalent COVID-19 booster vaccination in adults with and without prior exposure to SARS-CoV-2


As severe acute respiratory coronavirus 2 (SARS-CoV-2) variants continue to emerge, it is important to characterize immune responses against variants which can inform on protection efficacies following booster vaccination. In this study, neutralizing breadth and antigen-specific CD8⁺ T cell responses were analyzed in both infection-naïve and infection-experienced individuals following administration of a booster bivalent Wuhan-Hu-1+BA.4/5 Comirnaty® mRNA vaccine. Significantly higher neutralizing titers were found after this vaccination compared to the pre-third booster vaccination time point. Further, neutralizing breadth to omicron variants, including BA.1, BA.2, BA.5, BQ.1 and XBB.1, was found to be boosted following bivalent vaccination. SARS-CoV-2-specific CD8⁺ T cells were identified, but with no evidence that frequencies were increased following booster vaccinations. Spike protein-specific CD8⁺ T cells were the only responses detected after vaccination and non-spike-specific CD8⁺ T cells were only detected after infection. Both spike-specific and non-spike-specific CD8⁺ T cells were found at much lower frequencies than CD8⁺ T cells specific to cytomegalovirus (CMV), Epstein-Barr virus (EBV) and influenza (Flu). Taken together, these results show that the bivalent Wuhan-Hu-1+BA.4/5 Comirnaty® mRNA vaccine boosted the breadth of neutralization to newer SARS-CoV-2 variants and that vaccination is able to induce spike protein-specific CD8⁺ T cell responses, which are maintained longitudinally.

KEYWORDS
SARS-CoV-2, COVID-19, vaccination, neutralization, CD8 T cells
1 Introduction

Coronavirus disease 2019 (COVID-19), caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), has led to the deaths of millions globally (1). Following the introduction of approved COVID-19 vaccinations at the end of 2020, although protection from infection was not achieved, cases of severe COVID-19, hospitalization and death dropped significantly (1, 2). However, the protection efficacy of the original vaccines were found to diminish due to a combination of waning immunity and the emergence of new SARS-CoV-2 variants capable of evading prior-established immune responses (1, 3, 4). The evolving SARS-CoV-2 variants are generally characterized by mutations in the surface located spike (S) protein, which is the major protein for host cell entry and the major target for neutralizing antibody (nAb) responses. To improve immunity against emerging SARS-CoV-2 variants, administrations of booster vaccinations have been implemented, and vaccine formulas were updated from a monovalent ancestral S protein (Wuhan-Hu-1 variant) to a bivalent formula containing both the ancestral S protein (Wuhan-Hu-1 S mRNA) and an S protein from a more recent variant (omicron BA.1 or omicron BA.4/BA.5 S mRNA). This vaccine regimen was approved by the food and drug administration (FDA), United Kingdom (UK) government, European Medicines Agency (EMA) and therapeutic goods administration (TGA) for use in the United States (US), Canada, UK, Europe, and Australia. The three major vaccine regimes that have been approved include Comirnaty®, Spikevax® and Novavax®. However, as implementation of these updated vaccine formulas has been relatively slow, newer omicron variants emerged, such as omicron BQ.1 and omicron XBR.1, which have been shown to be highly resistant to neutralizing activity from both therapeutic monoclonal antibodies and sera from convalescent/vaccinated individuals (5–7). Recent studies have shown that nAb titers correlate with protection from developing severe COVID-19 (8, 9), thus, measurement of nAb titers following vaccination and/or infection is of high importance for prediction of protection efficacies.

Like nAb responses, recent evidence has suggested that SARS-CoV-2-specific CD8+ T cell responses correlate with protection from developing severe COVID-19 (10). Unlike nAb epitopes, which are greatly affected by the S protein changes found among the different SARS-CoV-2 variants, CD8+ T cell epitopes have been found to be more conserved (11, 12). However, while some CD8+ T cell epitopes have been proposed to be more immunodominant (12, 13), a large degree of heterogeneity has been found amongst both convalescent and vaccinated individuals. This is mostly due to the heterogeneity of class I major histocompatibility complexes (MHCs), which are responsible for presenting epitopes to CD8+ T cells. Each different type of MHC class I molecule is restricted to the type of epitopes it can present, which are linear amino acid sequences usually 8-11 amino acids in length. Using pools of CD8+ T cell epitopes, recent studies have shown that S protein-specific CD8+ T cells can be induced following COVID-19 vaccination (14–17), albeit to relatively low frequencies. However, it is not completely clear how these frequencies fluctuate with additional antigen exposures (infection or additional vaccination) nor have other studies compared the frequencies of SARS-CoV-2-specific CD8+ T cells to other antigen-specific CD8+ T cells or negative controls.

In the present study, the breadth of nAb responses in plasma were assessed in individuals with a COVID-19 vaccination history, including some with subsequent SARS-CoV-2 infection, with a particular focus on nAb responses after a fourth Pfizer/BioNTech (Comirnaty®) vaccine dose containing the ancestral (Wuhan-Hu-1) and BA.4/BA.5 bivalent formula. In addition, SARS-CoV-2-specific CD8+ T cells were analyzed longitudinally at pre-vaccination (and pre-infection) and approximately 1-month (1M) post vaccination for each vaccination (4 doses total). SARS-CoV-2-specific CD8+ T cells following vaccinations were split into S protein-specific and non-S protein-specific and compared between infection-naive individuals and infection-experienced individuals. Lastly, the frequencies of SARS-CoV-2-specific CD8+ T cells were compared to a non-specific epitope and to immunodominant epitopes found in cytomegalovirus (CMV), Epstein-Barr virus (EBV) and influenza (Flu).

2 Materials and methods

2.1 Study cohort

Participants for this study were selected from the clinical, virological and immunological COVID-19 (CVIC) study, which is a prospective cohort of individuals either infected by SARS-CoV-2 or vaccinated against COVID-19, as previously described (18–20). Participants for this study were selected based on whether they had received the bivalent (Wuhan-Hu-1+omicron BA.4/5 S mRNA) Comirnaty® booster vaccine. Any participants that received a Vaxzevria® or Spikevax® vaccination were excluded from this study. Thus, participants in this study received only Comirnaty® vaccinations. Longitudinal follow up included blood collection at pre-vaccination, 1M post-booster vaccination (second dose), 1M-post second booster vaccination (third dose), pre-third booster vaccination and 1M-post third booster vaccination (fourth dose).

2.2 Plasma and peripheral blood mononuclear cell isolation

Following blood collection in ethylenediaminetetraacetic acid (EDTA) tubes, blood was centrifuged at 1800 x g for 10 min with the brake turned off. Plasma was then collected and stored at -80°C in cryovials until use. The buffy coat containing peripheral blood mononuclear cells (PBMCs) was collected, combined with phosphate buffered saline (PBS) and added to Sepmate tubes (STEMcell Technologies) containing Ficoll (Sigma-Arich). The tubes were then centrifuged at 1200 x g for 10 min with the brake turned off. The supernatant was then poured into a fresh tube and topped up with PBS. The tubes were then centrifuged at 400 x g for 6 min and washed once more with PBS. Following this, the cells were resuspended in RPMI media (Gibco) and counted using a Countess II (Thermo Fisher Scientific). An equal part of pre-chilled freezing media (50% RPMI, 30% FBS and 20% dimethyl sulfoxide) was then added and cells were
frozen at -80°C in a CoolCell® container (Corning) in cryovials containing a minimum of 5 x 10^6 cells/vial. For long-term storage, the cryovials were moved to -150°C.

2.3 Screening for SARS-CoV-2 infection

As reported previously (19), to determine if a participant had been infected, plasma was screened longitudinally for the presence of anti-nucleocapsid (N) antibodies using a EuroImmun semi-quantitative enzyme-linked immunosorbent assay (ELISA) (PerkinElmer, cat#: El 2606-9601-2 G). All time points were compared to the pre-vaccination time point and a signal/noise ratio of 3.0 or more was considered positive. In addition, participants were required to report on testing positive for SARS-CoV-2 infection via routine diagnostic polymerase chain reaction (PCR) from nasopharyngeal swabs. Subsequently, any participants that became PCR positive had their nasopharyngeal swab sent to the Department of Microbiology, Copenhagen University Hospital, Hvidovre, Denmark, to see if a viral sequence could be recovered. RNA extraction of samples from swabs was done using PentaBase viral nucleic acid purification kit (PentaBase A/S). Sequencing libraries were prepared using MIDNIGHT-ONT/V3 primers and Rapid Barcode Sequencing Kit (Oxford Nanopore Technologies) following the manufacturer’s instructions. Sequencing was performed using R9.4.1 flow cells (Oxford Nanopore Technologies) on a GridION (Oxford Nanopore Technologies) using default parameters and high-accuracy base-calling. Reads were processed using MinKNOWs internal epiprome workflow (v.0.3.14) for SARS-CoV-2 with default parameters. Inside the workflow, variants were called using Nextclade (v.1.11.0) and Pangolin (v.4.0.5). All infections that had a variant identified can be found in Table 1.

2.4 Ethics

The study was approved by the Regional Ethical Committee (H-20025872) and Data Protection Agency (P-2020-3571), respectively, and was conducted in compliance with the Declaration of Helsinki guidelines. All individuals included in this study were 18 years or older and able to read and speak adequate Danish to provide written informed consent. Participants were required to self-report their age, sex, and ethnicity upon enrolment. Study data was collected and managed using research electronic data capture (REDCap) tools hosted at Copenhagen University Hospital, Hvidovre (21).

2.5 Neutralization assay

Initially, all participants included in this study were screened for plasma neutralization using a D614G SARS-CoV-2 isolate (DK-AHH1, Genbank accession number MZ049597), as described previously (19, 20, 22). In brief, virus was added to 2-fold serially diluted plasma at a 1:1 ratio and incubated at room temperature. Following 1h incubation, plasma/virus and antibody/virus complexes were then added to Vero E6 cells (RRID: CVCL_0574) seeded the day before (10^6 cells/well) in quadruplicate. After 48 hours incubation, the cells were fixed and stained. Spots representing virus infected cells were counted and single outliers were removed as previously described (23). The percentage neutralization was calculated as:

\[
\% \text{ Neutralisation} = 1 - \left( \frac{\text{Spot count}}{\text{Average spot count (virus only and healthy controls)}} \right) \times 100
\]

Following this initial screening, selected participants were then screened for plasma neutralization to other SARS-CoV-2 isolates including a delta isolate (DK-AHH3, accession number OP271297) (19), an omicron BA.1 isolate (DK-AHH4, accession number OP271296) (19), an omicron BA.2 isolate (DK-AHH5, accession number OP722493) (24), an omicron BA.5 isolate (DK-AHH6, accession number OP722492) (24), an omicron XBB.1.4 isolate (DK-AHH7, accession number OQ843560, described here for the first time) and an omicron BQ.1.1 isolate (DK-AHH8, clade 22E, accession number OQ843561, described here for the first time). All these isolates were grown in Vero E6 cell culture to obtain a viral stock. Viral stocks were then titrated to a multiplicity of infection (MOI) that could completely infect 10^4 Vero E6 cells without causing cytopathic effect. All neutralization experiments for these isolates were done as described above for DK-AHH1.

2.6 HLA typing

Human leukocyte antigen (HLA) typing of subjects was done using a protocol previously developed by others (25). In brief, subject DNA was obtained by lysing one vial of PBMCs using a DNeasy Blood & Tissue kit (Qiagen) according to the manufacturer’s instructions. This DNA was used as a template for PCR amplification of the HLA A allele using the primers and PCR cycling conditions previously described (25). Following PCR, the products were checked via gel agarose electrophoresis and purified using AMPure XP beads (Qiagen) according to the manufacturer’s instructions. Purified products were then pooled, and library preparation was conducted using a NEBNext Ultra II DNA Library Preparation kit (New England Biolabs). Next generation sequencing (NGS) was performed using the MiSeq platform (Illumina). The data was analyzed using Hisat-genotype analysis pipeline, as previously described (26).

2.7 Flow cytometry for screening of antigen-specific CD8+ T cells

Frozen PBMCs were thawed at room temperature and immediately transferred to PBS supplemented with 5% fetal bovine serum (FBS). The cell suspension was centrifuged at 500 x g for 5 min and the cell pellet was resuspended in PBS supplemented with 5% FBS. Cells were then stained in the dark with Fixable Viability Stain 780 (FVS780, BD Biosciences) for 20 min. The cells were then washed twice with PBS supplemented with 5% FBS and
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| Median (IQR) | 49 (46-57) | 74% female | Mean (SD) | 36 (7) | Mean (SD) | 215 (36) | Mean (SD) | 39 (7) |

DNS, did not show; N/A, not applicable. *N ELISA negative at prior time points.
resuspended in Brilliant Stain Buffer (BD Biosciences). The cells were split into four aliquots, one for the non-specific negative control Dextramer reagents, one for the CMV, EBV and Flu (CEF) Dextramer reagents, one for the SARS-CoV-2 S protein-specific Dextramer reagents and one for the SARS-CoV-2 non-S protein-specific Dextramer reagents. Dextramer staining was done in the dark for 30 min according to the manufacturer’s instructions. All Dextramer reagents were labelled with the PE fluorophore and the panel of Dextramer reagents can be found in Supplementary Table 1. These epitopes were selected from internal and published knowledge on immunodominant epitopes in relation to these specific HLA types (12, 13). Following this, the cells were stained in the dark for 20 min with anti-CD3-AF700 (BD Biosciences), anti-CD14-APC (BD Biosciences), anti-CD19-APC (BD Biosciences), anti-CD4-FLU395 (BD Biosciences) and anti-CD8-FITC (BD Biosciences). The cells were then washed three times with PBS supplemented with 5% FBS and resuspended in PBS supplemented with 5% FBS. Flow cytometry was then performed on a BD LSR Fortessa (5 laser) analyzer (BD Biosciences) using BD FACSDiva™ software version 6.1.3 (BD Biosciences). Downstream analysis of the flow cytometry data was done using FlowJo software version 10.8.1 (BD Biosciences).

2.8 Statistics

50% inhibitory dilution neutralization titers (ID_{50}) of plasma were calculated in GraphPad Prism (version 9.5.1). Each specific statistical test performed is indicated in the text and figure legends. In brief, data was checked for normal distribution by using QQ-plots and assessed using the Shapiro-Wilk test and Kolmogorov-Smirnov tests. Data that was not found to be normally distributed was analyzed using non-parametric tests and corrected for multiple comparisons as indicated. Data that did not pass the normal distribution tests were plotted with the median and 95% confidence interval. All analyses were two tailed and statistical significance was defined as a p value less than 0.05.

2.9 Role of funders

The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript. All authors had full access to all the data in the study and had final responsibility for the decision to submit for publication.

3 Results

3.1 Study participants vaccinated against COVID-19

A total of 109 participants vaccinated with a prime-boost Comirnaty® regimen (Wuhan-Hu-1 S mRNA-mRNA) were enrolled in the CVIC study and followed prospectively (18, 19). Among these individuals, 38 received both a third Wuhan-Hu-1 S mRNA Comirnaty® vaccination and a fourth bivalent (Wuhan-Hu-1+BA.4/5 S mRNA) Comirnaty® vaccination and were included in this study. Blood samples collected at pre-vaccination, 1M post-booster vaccination (second dose), 1M post-second booster vaccination (third dose), pre-third booster vaccination and 1M post-third booster vaccination (fourth dose) were analyzed (Figure 1A). Twenty-seven of these individuals had had a SARS-CoV-2 infection identified through N protein ELISA, of which 21 were confirmed via routine PCR testing. Twenty five of the 27 had an infection identified between their second booster and third booster Comirnaty® vaccinations and the remaining 2 had an identified infection between their booster and second booster Comirnaty® vaccinations (Table 1). Of the 21 with PCR confirmed SARS-CoV-2 infection, 9 had the infecting SARS-CoV-2 variant identified, of which 2 were omicron BA.1, 5 were BA.2, 1 was BA.5 and 1 was the alpha variant (Table 1). Regardless of when they were infected, the 27 individuals included in this study with an identified SARS-CoV-2 infection were termed “hybrid” and the remaining 11 that did not have a SARS-CoV-2 infection identified were termed “naïve”. The median age in the hybrid and naïve groups were 49 (interquartile range [IQR]=47-57) and 53 (IQR=48-58), respectively. The percentage of females was 74% in the hybrid group and 82% in the naïve group. A summary of participant information for the hybrid and naïve groups can be found in Table 1 and Table 2, respectively.

3.2 Assessment of neutralizing titers against an ancestral D614G SARS-CoV-2 isolate following serial vaccinations

Firstly, to assess the change in plasma neutralizing titers from the third booster vaccination dose containing the bivalent Wuhan-Hu-1+BA.4/5 S Comirnaty® mRNA, neutralizing titers against an ancestral D614G isolate were assessed in infectious culture systems (20, 22) at pre-third booster vaccination and 1M post-third booster vaccination time points in both naïve and hybrid groups (Figure 1B). Neutralization titers (50% inhibitory dilutions [ID_{50}]s) were determined by measuring viral infectivity following incubation with serially diluted plasma and comparing to the infectivity at the pre-vaccination time point at a single plasma dilution (1/10). For both the naïve and hybrid groups, significantly higher neutralizing titers were found at the 1M post-third booster vaccination time point when compared to their respective pre-third booster vaccination time point (p<0.001, Wilcoxon t tests). Compared to the naïve group at 1M post-third booster vaccination, the hybrid group had significantly higher neutralizing titers (p=0.0175, Mann-Whitney U test). Next, neutralizing titers were assessed longitudinally in each group at 1M post-booster, 1M post-second booster and 1M post-third booster vaccination time points (Figure 1C). Although both groups appeared to have the highest neutralizing titers against the ancestral D614G isolate at the 1M post-second booster vaccination time point, there were no significant differences found in either group between the time points (p>0.05, Kruskal-Wallis tests).
Study cohort summary and examination of neutralizing titers against the ancestral D614G SARS-CoV-2 isolate in the naïve (n=11) and hybrid (n=27) groups following vaccinations against COVID-19. (A) Of the 109 participants in the CVIC study cohort, 38 had longitudinal follow-up that included blood collection at 1-month (1M) after bivalent (Wuhan-Hu-1+BA.4/5 S mRNA) third booster vaccination. Collection dates (blood vials) were done at pre-vaccination, 1M post-booster vaccination, 1M post-second booster vaccination, pre-third booster vaccination and 1M post-third booster vaccination. The blue syringes represent vaccination with the monovalent Comirnaty® ancestral formula (Wuhan-Hu-1 S mRNA) and the yellow syringe represents vaccination with the Comirnaty® bivalent (Wuhan-Hu-1+BA.4/5 S mRNA). Eleven of the 38 included did not have a SARS-CoV-2 infection detected throughout follow-up (blue line, termed “naïve”). Twenty seven of the 38 were found to have been infected with SARS-CoV-2 (orange line, termed “hybrid”). Twenty five of the 27 had an infection detected between their second booster and third booster Comirnaty® vaccinations and two had an infection detected between their booster and second booster Comirnaty® vaccinations. (B) Comparison of neutralizing titers against the ancestral D614G SARS-CoV-2 isolate between the pre-third booster and post-third booster vaccination time points in the naïve (blue and purple) and hybrid (orange and red) groups. Neutralizing titers were found to be significantly boosted following third booster vaccination in both the naïve (p=0.0010, Wilcoxon t test) and hybrid (p<0.0001, Wilcoxon t test) groups. When the two groups were compared at the 1M post-third booster vaccination time point (right graph), the hybrid group was found to have significantly higher neutralizing titers (p=0.0175, Mann-Whitney U test). (C) Longitudinal neutralizing titers at 1M post-booster (post-boost), 1M post-second booster (post-2nd boost) and 1M post-third booster (post-3rd boost) vaccinations in the naïve group (blue) and hybrid group (orange). *p<0.05, ***p<0.001 and ****p<0.0001.
3.3 Assessment of neutralizing breadth against delta and omicron SARS-CoV-2 isolates after second and third booster vaccinations

To assess neutralizing breadth in the mRNA vaccination groups with and without prior SARS-CoV-2 infection, 11 participants from the hybrid group were selected to match the 11 participants in the naïve group to allow a more direct comparison between the two groups. The participants from the hybrid group were selected based on matched age, sex, complete longitudinal follow up, and only having a detected SARS-CoV-2 infection between their post-second booster and pre-third booster vaccination time points (representing a time when the SARS-CoV-2 omicron BA.2 variant was dominant in Denmark). A summary of the 11 selected participants from the hybrid group can be found in Supplementary Table 2. Neutralizing breadth was measured against a panel of SARS-CoV-2 isolates representing delta and omicron (BA.1, BA.2, BA.5, BQ.1.1 and XBB.1.4) variants. Firstly, neutralizing breadth was compared at 1M post-second booster and 1M post-third booster vaccination time points within the naïve (Figure 2A) group and at 1M post-second booster, pre-third booster (post-infection) and 1M post-third booster vaccination time points in the hybrid group (Figure 2B). For the naïve group, when compared to the 1M post-second booster vaccination time point, significantly higher neutralizing titers were observed at the 1M post-third booster vaccination time point (p<0.01, Bonferroni-Dunn corrected Wilcoxon t tests). Furthermore, at the pre-third booster vaccination time point, neutralizing titers were significantly higher against the omicron BA.1, BA.2 and BA.5 isolates when compared to the post-second booster vaccination time point (p<0.05, Bonferroni-Dunn corrected Wilcoxon t tests). When the pre-third booster and post-third booster vaccination time points were compared, significantly higher neutralizing titers were detected at the post-third booster vaccination time point against all tested SARS-CoV-2 isolates (p<0.05, Bonferroni-Dunn corrected Wilcoxon t tests). To see if the breadth of neutralization in the hybrid group was superior to the naïve group, neutralizing titers were compared between these two groups at 1M post-second booster vaccination (Figure 2C) and at 1M post-third booster vaccination (Figure 2D). At 1M post-second booster vaccination, no observable differences were seen between the naïve and hybrid groups. At the 1M post-third booster vaccination time point, although the hybrid group had noticeably higher neutralizing titers against all the isolates tested, significance was only detected against the delta, omicron BA.1 and omicron BA.2 isolates (p<0.05, Bonferroni-Dunn corrected Mann-Whitney U tests).

3.4 Detection of antigen-specific CD8+ T cell responses

In order to identify SARS-CoV-2-specific CD8+ T cell responses, we selected S protein-specific and non-S protein-specific SARS-CoV-2 epitopes displayed by MHC on PE-labelled Dextramer reagents specific to HLA types A*01:01, A*02:01 and

### Table 2: Summary of participant information for the naïve group vaccinated against SARS-CoV-2.

<table>
<thead>
<tr>
<th>Patient ID</th>
<th>Age (years)</th>
<th>Sex</th>
<th>Ethnicity</th>
<th>Second booster vaccination (date)</th>
<th>Second booster vaccination time point (days post vaccination)</th>
<th>Third booster vaccination (date)</th>
<th>Third booster vaccination time point (days post vaccination)</th>
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<td>35</td>
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<td>5 10 2022</td>
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</tr>
<tr>
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<td>2 11 2022</td>
<td>33</td>
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<tr>
<td>mRNA-102</td>
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<td>22 10 2021</td>
<td>DNS</td>
<td>5 10 2022</td>
<td>49</td>
</tr>
</tbody>
</table>

Median (IQR) 53 (48-58) 18% male

82% female  Mean (SD) 38 (9)  Mean (SD) 39 (6)

DNS, did not show.
Next, the 11 participants from the naïve group and the 11 matched participants from the hybrid group (i.e., the same 11 participants as studied above for cross-neutralizing antibodies) were HLA typed to identify those with suitable HLA A alleles. Of these participants, 10/11 of the naïve participants and 9/11 of the hybrid participants were found to have matching HLA A alleles (Supplementary Table 3). The MHC class I Dextramer reagents were used to identify SARS-CoV-2-specific CD8+ T cells in PBMCs isolated from peripheral blood at the same time points used in the neutralization analyses (Figure 1A). For each time point, analysis was done using four pools of Dextramer reagents, one containing negative control Dextramer reagents, one containing Dextramer reagents specific for other viral epitopes (CMV, EBV and Flu), one containing SARS-CoV-2 S protein-specific Dextramer reagents and one containing SARS-CoV-2 non-S protein-specific Dextramer reagents. Each pool of Dextramer reagents was analyzed separately from one another using flow cytometry and gated as shown in Figure 3. Any time points that had <5,000 CD8+ T cells were excluded from further analyses. In the naïve group, three showed consistently low levels of CD8+ T cells (<5,000 CD8+ T cells) and were excluded, meaning that only 7 subjects were included in the naïve group. A summary of the time points included and the detected antigen-specific CD8+ T cell frequencies can be found in Supplementary Table 4.

3.5 Assessment of S protein-specific CD8+ T cells

Given that the vaccine formulas used in this study only contained the SARS-CoV-2 S protein, CD8+ T cell frequencies were first measured against epitopes specific to the SARS-CoV-2 S protein (Figure 4). For all individuals, S protein-specific CD8+ T cells were not detected at the pre-vaccination time point.
In two individuals, S protein-specific CD8+ T cells were not detected at any time point. For the remaining individuals, all follow-up time points showed continuously detectable S protein-specific CD8+ T cells at varying frequencies (Figure 4A). However, while some subjects showed a trend for increasing S protein-specific CD8+ T cell frequencies after each antigen exposure (i.e., infection or booster vaccination), no significant differences in the frequencies of S protein-specific CD8+ T cells were found at any time point (excluding comparisons to the pre-vaccination time point, p>0.05, Dunn’s corrected Kruskal-Wallis test). To see if there were any differences in the frequencies of S protein-specific CD8+ T cells between the naïve and hybrid groups, these groups were separated and compared at each time point (Figure 4B). Although slightly higher frequencies of S protein-specific CD8+ T cells were detected in the hybrid group at the post-third booster vaccination time point, no significant differences were detected at any time point (p>0.05, Holm Sidak corrected multiple Mann-Whitney U tests).

(Figure 4A). In two individuals, S protein-specific CD8+ T cells were not detected at any time point. For the remaining individuals, all follow-up time points showed continuously detectable S protein-specific CD8+ T cells at varying frequencies (Figure 4A). However, while some subjects showed a trend for increasing S protein-specific CD8+ T cell frequencies after each antigen exposure (i.e., infection or booster vaccination), no significant differences in the frequencies of S protein-specific CD8+ T cells were found at any time point (excluding comparisons to the pre-vaccination time point, p>0.05, Dunn’s corrected Kruskal-Wallis test). To see if there were any differences in the frequencies of S protein-specific CD8+ T cells between the naïve and hybrid groups, these groups were separated and compared at each time point (Figure 4B). Although slightly higher frequencies of S protein-specific CD8+ T cells were detected in the hybrid group at the post-third booster vaccination time point, no significant differences were detected at any time point (p>0.05, Holm Sidak corrected multiple Mann-Whitney U tests).

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3.6 Assessment of non-S protein-specific and CEF-specific CD8+ T cells

Given that some of the individuals became infected, immunodominant epitopes outside of the S protein (i.e., non-S protein-specific epitopes) were assessed longitudinally (Figure 5A). Detection of non-S protein-specific CD8+ T cells only occurred at the pre-third booster vaccination and post-third booster vaccination time points (i.e., post-infection time points, Figure 5A). When these two time points were compared, no significant differences were found (p>0.05, Mann-Whitney U test). Detection of these CD8+ T cells only occurred in individuals in the hybrid group (Figure 5B). When compared at the post-third booster vaccination time point, the hybrid group had significantly higher non-S protein-specific CD8+ T cell frequencies than the naïve group (p=0.005, Mann-Whitney U test, Figure 5B). No other time points were compared as non-S protein-specific CD8+ T cells were not detected.

Longitudinal assessment of CEF-specific CD8+ T cells showed that they could be detected at all time points for all individuals at varying frequencies (Figure 5C). However, no significant differences were detected between any of the time points (p>0.05, Dunn’s corrected Kruskal-Wallis test). Similarly, when the frequencies of CEF-specific CD8+ T cells were compared between the naïve and hybrid groups, no significant differences were found (p>0.05, Holm-Sidak corrected multiple Mann-Whitney U tests, Figure 5D).

4 Discussion

Since the release of several COVID-19 vaccination strategies at the end of 2020, there has been a race between the emergence of SARS-CoV-2 variants and vaccination capable of protecting against evolving variants. With recent evidence showing that the newer BQ and XBB omicron variants are highly resistant to previously established neutralizing responses (5, 6), it’s important for vaccine formulas to be updated to help induce immunity that can protect against current circulating variants. While updated COVID-19 vaccine formulas have been shown to induce improved breadth of
neutralization to emerging variants (5, 6), the level of neutralization is still significantly lower than that against the ancestral SARS-CoV-2 variant (Wuhan-Hu-1). However, continued antigen exposure, whether that be through vaccination or through infection, has shown improved breadth of neutralization to multiple SARS-CoV-2 variants (19). Similarly, the results presented in this study support the continued development of neutralizing breadth through additional antigen exposures, whether that be from additional vaccination, infection, or both. Particularly, individuals who had received the new Comirnaty® bivalent vaccine (third booster vaccination) showed a large improvement in neutralization titers to the omicron variants tested. Although limited, other studies that have assessed the bivalent Spikevax® vaccine (mRNA-1273.214) from Moderna have also shown comparable results to that seen in this study (27, 28). That is, the bivalent booster vaccination increased breadth of neutralization to omicron variants with a greater boost seen in those with prior infection. Interestingly, however, neutralization to the ancestral D614G isolate in this study, although boosted compared to pre-bivalent vaccination, was not brought back to comparable levels to that seen at post-second booster vaccination, unless individuals had had an infection (hybrid group). In a previous report assessing neutralization in individuals that received two booster vaccinations and individuals that received three booster vaccinations, while the monovalent third booster vaccination showed substantially increased neutralization titers against the ancestral D614G coronavirus pseudoparticle (CoVpp) compared to the bivalent third booster vaccination, the bivalent third booster vaccination showed much higher neutralization titers against the ancestral D614G CoVpp when compared to the neutralization titers after two booster vaccinations (29). However, this prior study did not use the same individuals for assessing neutralizing titers after three and four vaccine doses and failed to time-match the samples based on sample collection following vaccination, which has been shown to be important as neutralizing titers rapidly wane after 1M post-vaccination (19, 30). Therefore, this may explain the differences with the results found in the present study. This would make sense given that it is likely that memory B cells would be competing for the vaccine-induced antigens, and that having a lower amount of previously encountered antigen (Wuhan-Hu-1 S protein) would limit memory B cells from being activated, thus forcing differentiation and maturation of naïve B cells to the newly encountered omicron S antigens (BA.4/5 S protein). Therefore, compared to the third dose vaccination, which is exclusively the Wuhan-Hu-1 S protein, the bivalent vaccination would not induce as high neutralizing titers to the ancestral D614G isolate. However, given that the ancestral D614G SARS-CoV-2 variant is not circulating anymore, improvement of neutralizing titers against the newer BQ and XBB omicron variants is of higher importance. The data presented here, which is supported by others (6, 27, 30), shows that Comirnaty® bivalent vaccination can indeed boost neutralization titers to the newer omicron variants. Although differences in monovalent and bivalent third booster vaccinations were not assessed in this study, others have shown that a bivalent third booster vaccination is superior at inducing improved neutralization titers against newer omicron variants (27, 31).

Although newer SARS-CoV-2 variants have been shown to be increasingly evasive to both infection- and vaccine-induced nAb responses, known CD8+ T cell epitopes have been found to be much more conserved between the different variants (11, 12). Given the apparent importance of CD8+ T cells in the complete clearance of SARS-CoV-2 infection (32, 33), antigen-specific CD8+ T cells were investigated in this study. Unlike previous studies assessing SARS-CoV-2-specific CD8+ T cell frequencies (13, 34, 35), this study compared the SARS-CoV-2-specific CD8+ T cell frequencies to the frequencies of other virus-specific CD8+ T cell responses (CEF pool). For SARS-CoV-2, the assay used here allowed tracking of the virus-specific T cell dynamics in response to both vaccination and infection. It was clear to see that, following prime-boost vaccination, S protein-specific CD8+ T cells could be detected and, following infection, detection of non-S protein-specific CD8+ T cells was observed. Importantly, these antigen-specific CD8+ T cells were not detected at pre-vaccination or at pre-infection time points, indicating that the CD8+ T cell epitopes employed in this study were specific to SARS-CoV-2 and that these epitopes were successful at identifying antigen-specific CD8+ T cells corresponding to the exposed antigen. When compared to other studies looking at SARS-CoV-2-specific CD8+ T cells, the frequencies found in this study are similar to that reported by others (13, 34, 35). Virus-specific T cell responses directed against CMV, EBV and Flu were detected at much higher frequencies compared to the SARS-CoV-2-specific CD8+ T cells, which could be due to several reasons. Firstly, CMV, EBV and Flu specific T cells were detected as one group, thus the CEF-specific cell numbers observed is the sum of responses to 3 viruses and not just one as for the SARS-CoV-2 responses measured. Secondly, these CD8+ T cells were likely formed from an infection with higher antigenic exposure than a vaccine, allowing greater expansion of antigen-specific CD8+ T cells prior to antigen clearance.

Interestingly, when the antigen-specific CD8+ T cell frequencies were examined longitudinally to either SARS-CoV-2 or CEF, no fluctuation in cell numbers was observed, even after additional antigen exposure (i.e., SARS-CoV-2 booster vaccination or infection). This is comparable to what has been observed in another study examining SARS-CoV-2-specific CD8+ T cells longitudinally (17). Given that it has been shown that the CD8+ T cell response to vaccinations peak around 10 days post-antigen exposure (36, 37), it is possible that frequencies of antigen-specific CD8+ T cells detected in this study, and by others (17), are after the antigen-specific CD8+ T cells have contracted back to a plateau, meaning that the detected antigen-specific CD8+ T cell frequencies in this study may have been past the expansion phase and, therefore, likely to be in a memory state. This would make sense considering SARS-CoV-2-specific memory CD8+ T cells have been shown to be long-lasting (38).

Although the epitopes used in this study were carefully selected based on previous reports of these epitopes being found at high rates among the HLA types selected for this study (12, 13), inclusion of additional epitopes may help improve detection frequencies of the SARS-CoV-2 non-S protein-specific CD8+ T cell frequencies. However, even epitopes like TTDP5SLFGLRY, which has been reported to be highly immunodominant in HLA A*01:01 individuals (13), were not found at high frequencies in A*01:01
individuals that acquired a SARS-CoV-2 infection in this study. However, this could be due to the participants having already established S protein-specific CD8+ T cell responses from COVID-19 vaccination, which will then be favored upon antigen re-exposure, especially considering the degree of conservation of CD8+ T cell epitopes between the SARS-CoV-2 variants. This phenomenon has been reported in individuals vaccinated against COVID-19 and suggested to be an effect of ‘immune imprinting’ (39). However, there is no evidence to suggest that immune imprinting induces better or worse disease outcomes after infection.

Overall, this study supports that the administration of a bivalent (Wuhan-Hu-1+BA.4/5 S mRNA) Comirnaty® vaccination regime induces improved neutralizing antibody responses to omicron variants for both infection-naïve and infection-experienced individuals. Given the relationship between neutralizing titers and protection from severe COVID-19, the improved neutralizing responses seen from the bivalent Comirnaty® booster vaccination are likely to improve protection efficacies. Furthermore, although SARS-CoV-2-specific CD8+ T cells were not found to be boosted following serial vaccinations, the number of specific T cells were sustained in those that had detectable responses. Lastly, there was no evidence of infection-experienced individuals having superior S protein-specific CD8+ T cell responses to infection-naïve vaccinated individuals, suggesting that infection-induced responses are not superior to vaccine-induced responses.

Data availability statement

The original contributions presented in the study are included in the article/Supplementary Material. Further inquiries can be directed to the corresponding author.

Ethics statement

The studies involving humans were approved by the Regional Ethical Committee (H-20025872) and Data Protection Agency (P-2020-357). The studies were conducted in accordance with the local legislation and institutional requirements. The participants provided their written informed consent to participate in this study.

Author contributions

AU: Conceptualization, Data curation, Formal analysis, Funding acquisition, Investigation, Methodology, Software, Validation, Visualization, Writing – original draft, Writing – review & editing. CS: Funding acquisition, Resources, Writing – review & editing. KJ: Data curation, Formal analysis, Methodology, Resources, Validation, Writing – review & editing. AB: Investigation, Writing – review & editing. CF-A: Investigation, Writing – review & editing. LM: Investigation, Writing – review & editing. DI: Data curation, Methodology, Writing – review & editing. SLV: Resources, Writing – review & editing. JC: Formal analysis, Resources, Writing – review & editing, Investigation. MP: Resources, Writing – review & editing, Supervision. UF: Investigation, Methodology, Writing – review & editing. SR: Supervision, Writing – review & editing, Funding acquisition, Methodology, Resources. LB: Formal analysis, Resources, Supervision, Writing – review & editing. NW: Funding acquisition, Resources, Supervision, Writing – review & editing. JB: Conceptualization, Data curation, Formal analysis, Funding acquisition, Methodology, Project administration, Resources, Supervision, Validation, Writing – original draft, Writing – review & editing.

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Conflict of interest

Authors KJ, DI, and LB are employed by the company Immudex ApS.

The remaining authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fimmu.2024.1353353/full#supplementary-material

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